## CLAIMS

## What is claimed is:

|      | 1.  | A method of inhibiting amyloid plaque formation in a cell population     |
|------|-----|--|
|      |     | comprising contacting said cell population with an effective amount of a |
| 5    |     | compound selected from:  |
|      |     | (S) 6-(2-Imidazol-1-yl-1-phenyl-ethoxy)-5-phenethyl-3,4-dihydro          |
|      |     | 2H-naphthalen-1-one;   |
|      |     | 6-[2-(1H-Imidazol-4-yl)-ethoxy]-3,4-dihydro-2H-naphthalen-1-             |
|      |     | one;   |
| 10   |     | E-(+/-)6-(2-Imidazol-1-yl-1-phenyl-ethoxy)-2-thiophen-2-                 |
|      |     | ylmethylene-3,4-dihydro-2H-naphthalen-1-one;                             |
|      |     | 6-[1-(4-Chloro-phenyl)-2-imidazol-1-yl-ethoxy]-3,4-dihydro-2H-           |
|      |     | napththalen-1-one racemic;   |
|      |     | (R) 6-(2-Imidazol-1-yl-1-phenyl-ethoxy)-3,4-dihydro-2H-                  |
| 15   |     | naphthalen-1-onc;  |
|      |     | 6-(2-Imidazol-1-yl-1-phenyl-ethoxy)-4-phenyl-3,4-dihydro-2H-             |
|      |     | naphthalen-1-one racemic;  |
|      |     | 6-(2-Imidazol-1-yl-1-phenyl-ethoxy)-5-isopropoxymethyl-3,4-              |
|      | + 5 | dihydro-2H-naphthalen-1-one;   |
| 20   |     | (S) 6-(2-Imidazol-1-yl-1-phenyl-ethoxy)-5-phenylaminomethyl-             |
|      |     | 3,4-dihydro-2H-naphthalen-1-one;   |
|      |     | (S) 6-(2-Imidazol-1-yl-1-phenyl-ethoxy)-5-[2-(4-                         |
|      |     | fluorophenyl)ethyl]-3,4-dihydro-2H-naphthalen-1-one;                     |
|      |     | (S) 5-Benzenesulfonylmethyl-6-(2-imidazol-1-yl-1-phenyl-                 |
| 25 · | •   | ethoxy)-3,4-dihydro-2H-naphthalen-1-one;                                 |
|      |     | (S) 6-(2-Imidazol-1-yl-1-phenyl-ethylsulfanyl)-5-phenethyl-3,4-          |
|      |     | dihydro-2H-naphthalen-1-one;   |
|      |     | (S) 6-(2-Imidazol-1-yl-1-phenyl-ethoxy)-5-(2-pyridin-2-yl-ethyl)         |
| . ÷  |     | 3,4-dihydro-2H-naphthalen-1-one;   |
| 30   |     | 6-(2-Imidazol-1-yl-1-phenyl-ethoxy)-5-(2-pyridin-4-yl-ethyl)-3.4         |

dihydro-2H-naphthalen-1-one;

- 4-(5-Oxo-1-phenethyl-5,6,7,8-tetrahydro-naphthalen-2-yloxy)-4-phenyl-butyric acid;
- 6-[2-(3-Benzyl-3H-imidazol-4-yl)-ethoxy]-5-phenethyl-3,4-dihydro-2H-naphthalen-1-one; trifluoro-acetate;

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- (S) [1-{(4-Benzyloxy-benzyl)-[(2-methyl-2-phenyl-propylcarbamoyl)-methyl]-carbamoyl}-2-(3H-imidazol-4-yl)-ethyl]-carbamic acid benzyl ester;
- (S) [2-(1H-Imidazol-4-yl)-1-((4-methyl-benzyl)-{[(1-phenyl-cyclobutylmethyl)-carbamoyl]-methyl}-carbamoyl)-ethyl]-carbamic acid benzyl ester;
- 1-Methyl-4-(3-chlorophenyl)-6-[(4-chlorophenyl)-(1-methylimidazol-5-yl)aminomethyl]-2,3-dihydroquinolin-2-one; and
  (S) [1-{(4-Benzyloxy-benzyl)-[(2-benzyloxy-ethylcarbamoyl)-methyl]-carbamoyl}-2-(1H-imidazol-4-yl)ethyl]-carbamic acid benzyl ester.
- The method of Claim 1, wherein the compound administered is
   (S) 6-(2-Imidazol-1-yl-1-phenyl-ethoxy)-5-phenethyl-3,4-dihydro-2H-naphthalen-1-one.
- 3. The method of Claim 1, wherein said cell is in culture.
- 20 4. The method of Claim 1, wherein said cell population is in an animal.
  - The method of Claim 1, wherein said cell is a brain cell, a pancreatic cell, a kidney cell, a cardiac cell, a neuronal cell, or a thyroid cell.
  - A method for inhibiting amyloidosis in a patient comprising administering to said patient an amount effective to inhibit plaque formation of a compound selected from
    - (S) 6-(2-Imidazol-1-yl-1-phenyl-ethoxy)-5-phenethyl-3,4-dihydro-2H-naphthalen-1-one;

|    | <del></del>   |
|----|---|
|    | 6-[2-(1H-Imidazol-4-yl)-ethoxy]-3,4-dihydro-2H-naphthalen-1-      |
|    | one;  |
|    | E-(+/-)6-(2-Imidazol-l-yl-l-phenyl-ethoxy)-2-thiophen-2-          |
|    | ylmethylene-3,4-dihydro-2H-naphthalen-1-one;                      |
| 5  | 6-[1-(4-Chloro-phenyl)-2-imidazol-1-yl-ethoxy]-3,4-dihydro-2H-    |
|    | napththalen-1-one racemic;  |
|    | (R) 6-(2-Imidazol-1-yl-1-phenyl-ethoxy)-3,4-dihydro-2H-           |
|    | naphthalen-l-one;   |
|    | 6-(2-Imidazol-1-yl-1-phenyl-ethoxy)-4-phenyl-3,4-dihydro-2H-      |
| 10 | naphthalen-1-one racemic;   |
|    | 6-(2-Imidazol-1-yl-1-phenyl-ethoxy)-5-isopropoxymethyl-3,4-       |
|    | dihydro-2H-naphthalen-1-one;                                      |
| ı  | (S) 6-(2-Imidazol-1-yl-1-phenyl-ethoxy)-5-phenylaminomethyl-      |
|    | 3,4-dihydro-2H-naphthalen-1-one;                                  |
| 15 | (S) 6-(2-Imidazol-1-yl-1-phenyl-ethoxy)-5-[2-(4-                  |
|    | fluorophenyl)ethyl]-3,4-dihydro-2H-naphthalen-1-one;              |
|    | (S) 5-Benzenesulfonyimethyi-6-(2-imidazol-i-yl-i-phenyl-          |
|    | ethoxy)-3,4-dihydro-2H-naphthalen-1-one;                          |
|    | (S) 6-(2-Imidazol-1-yl-1-phenyl-ethylsulfanyl)-5-phenethyl-3,4-   |
| 20 | dihydro-2H-naphthalen-1-one;                                      |
|    | (S) 6-(2-Imidazol-1-yl-1-phenyl-ethoxy)-5-(2-pyridin-2-yl-ethyl)- |
|    | 3,4-dihydro-2H-naphthalen-1-one;                                  |
|    | 6-(2-Imidazol-1-yl-1-phenyl-ethoxy)-5-(2-pyridin-4-yl-ethyl)-3,4- |
|    | dihydro-2H-naphthalen-1-one;                                      |
| 25 | 4-(5-Oxo-1-phenethyl-5,6,7,8-tetrahydro-naphthalen-2-yloxy)-4-    |
|    | phenyl-butyric acid;  |
|    | 6-[2-(3-Benzyl-3H-imidazol-4-yl)-ethoxy]-5-phenethyl-3,4-         |
|    | dihydro-2H-naphthalen-1-one; trifluoro-acetate;                   |
|    | (S) [1-{(4-Benzyloxy-benzyl)-[(2-methyl-2-phenyl-                 |
| 30 | propylcarbamoyl)-methyl]-carbamoyl}-2-(3H-imidazol-4-yl)-ethyl]-  |
|    | carbamic acid benzyl ester;                                       |
|    |   |

(S) [2-(1H-Imidazol-4-yl)-1-((4-methyl-benzyl)-{{(1-phenyl-cyclobutylmethyl)-carbamoyl}-methyl}-carbamoyl)-ethyl]-carbamic acid benzyl ester;

l-Methyl-4-(3-chlorophenyl)-6-[(4-chlorophenyl)-(1-methylimidazol-5-yl)aminomethyl]-2,3-dihydroquinolin-2-one; and
(S) [1-{(4-Benzyloxy-benzyl)-[(2-benzyloxy-ethylcarbamoyl)-methyl]-carbamoyl}-2-(1H-imidazol-4-yl)ethyl]-carbamic acid benzyl ester.

7. The method of Claim 6, wherein said amyloidosis is associated with Alzheimer's disease.

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- 8. A method of treating Alzheimer's disease comprising administering to a patient in need of treatment an effective amount of
  - (S) 6-(2-Imidazol-1-yl-1-phenyl-ethoxy)-5-phenethyl-3,4-dihydro-2H-naphthalen-1-one.